

13/12/2006,10521325b.trn

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PASSWORD:

* * * * * RECONNECTED TO STN INTERNATIONAL * * * * *
SESSION RESUMED IN FILE 'HCAPLUS' AT 14:06:29 ON 07 DEC 2006
FILE 'HCAPLUS' ENTERED AT 14:06:29 ON 07 DEC 2006
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COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	50.80	655.24

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-3.00	-41.25

=> file reg

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	50.80	655.24

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-3.00	-41.25

FILE 'REGISTRY' ENTERED AT 14:06:41 ON 07 DEC 2006
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Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 6 DEC 2006 HIGHEST RN 914980-83-9
DICTIONARY FILE UPDATES: 6 DEC 2006 HIGHEST RN 914980-83-9

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH June 30, 2006

Please note that search-term pricing does apply when conducting SmartSELECT searches.

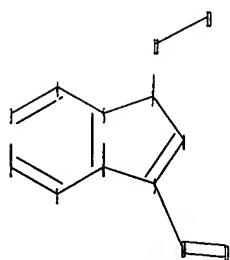
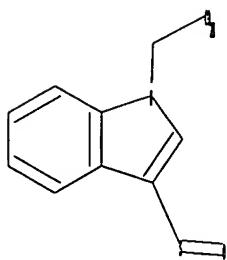
REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

<http://www.cas.org/ONLINE/UG/regprops.html>

=>
Uploading C:\Program Files\Stnexp\Queries\105121325.str

Young, Shawquia, Page 1

13/12/2006, 10521325b.trn



chain nodes :

10 11 12 13

ring nodes :

1 2 3 4 5 6 7 8 9

chain bonds :

7-12 9-10 10-11 12-13

ring bonds :

1-2 1-6 2-3 3-4 4-5 5-6 5-7 6-9 7-8 8-9

exact/norm bonds :

5-7 6-9 7-8 7-12 8-9 9-10 10-11

exact bonds :

12-13

normalized bonds :

1-2 1-6 2-3 3-4 4-5 5-6

Match level :

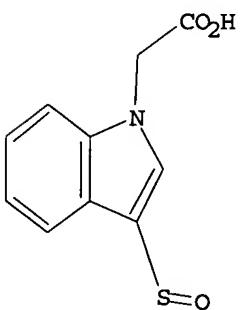
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11:CLASS 12:CLASS 13:CLASS

L10 STRUCTURE UPLOADED

=> d 110

L10 HAS NO ANSWERS

L10 STR



Structure attributes must be viewed using STN Express query preparation.

=> s 110

SAMPLE SEARCH INITIATED 14:07:11 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 7 TO ITERATE

13/12/2006, 10521325b.trn

100.0% PROCESSED 7 ITERATIONS 6 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 7 TO 298
PROJECTED ANSWERS: 6 TO 266

L11 6 SEA SSS SAM L10

=> s l10 full
FULL SEARCH INITIATED 14:07:15 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 105 TO ITERATE

100.0% PROCESSED 105 ITERATIONS 58 ANSWERS
SEARCH TIME: 00.00.01

L12 58 SEA SSS FUL L10

=> file hcplus			
COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION	
FULL ESTIMATED COST	166.94	822.18	
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION	
CA SUBSCRIBER PRICE	0.00	-41.25	

FILE 'HCAPLUS' ENTERED AT 14:07:19 ON 07 DEC 2006
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FILE COVERS 1907 - 7 Dec 2006 VOL 145 ISS 24
FILE LAST UPDATED: 6 Dec 2006 (20061206/ED)

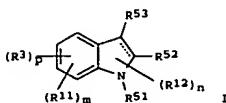
New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

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L13 4 L12

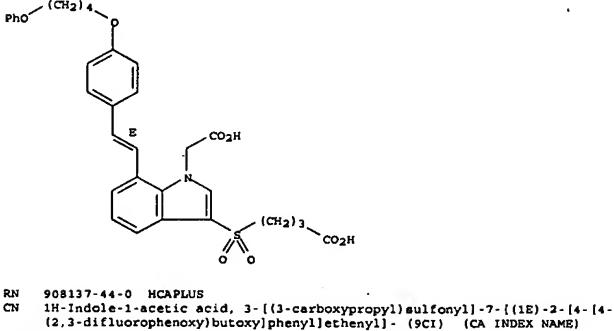
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L13 ANSWER 1 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN
 ED Entered STN: 31 Aug 2006
 GI



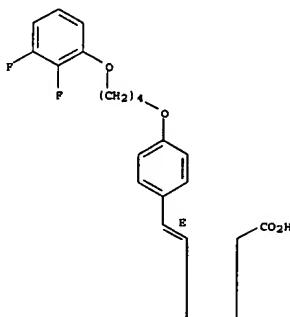
AB Indole compds. represented by the general formula (I) or salts or solvates thereof or prodrugs thereof [R11, R12 = substituents; two of R51, R52, and R53 are independently groups having a (un)protected acidic group and the remaining one group is H or a substituent; R3 = a substituent; m = an integer of 0-4; n = an integer of 0-2; p = 0, 1] are prepared. These compds. have a leukotriene receptor antagonistic effect and are expected to be more effective than those of the leukotriene receptor antagonists currently used in clin. medicine. They are therefore useful as agents for prevention and/or treatment of a leukotriene-mediated disease such as a respiratory disease, e.g., bronchial asthma, chronic obstructive pulmonary disease, pulmonary emphysema, chronic bronchitis, pneumonia (e.g., interstitial pneumonia), severe acute respiratory syndrome (SARS), acute respiratory distress syndrome (ARDS), allergic rhinitis, sinusitis (e.g., acute sinusitis, chronic sinusitis), and pulmonary fibrosis, and as expectorants or antitussives. Thus, Me 4-bromo-1-(4-methoxy-4-oxobutyl)-1H-indole-3-carboxylic acid was coupled with 4-vinylphenyl acetate in the presence of palladium acetate and tri(2-methylphenyl)phosphine in a solution of Et₃N in MeCN at 85° for 2 h to give Me 4-[{E}-2-[4-(acetoxyphenyl)ethenyl]-1-(4-methoxy-4-oxobutyl)-1H-indole-3-carboxylic acid. The latter compound was deacetylated by treatment with K₂CO₃ in a mixture of methanol and THF at room temperature for 2 h and etherified with 1-chloro-4-phenylbutane in the presence of NaI and K₂CO₃ in DMF at 95° for 2 h to give Me 4-[{E}-2-[4-(4-phenylbutoxy)phenyl]ethenyl]-1H-indole-3-carboxylic acid which was stirred with a mixture of 1 M aqueous NaOH solution, THF, and MeOH and acidified with 1-2 M aqueous HCl solution to give 1-(3-carboxypropyl)-4-[{E}-2-[4-(4-phenylbutoxy)phenyl]ethenyl]-1H-indole-3-carboxylic acid. 4-(1-(Carboxymethyl)-7-[{E}-2-[4-(4-phenoxybutoxy)phenyl]ethenyl]-1H-indol-3-yl)butanoic acid at 10 mg/kg p.o. in vivo inhibited the ovalbumin-induced constriction of airway in guinea pigs. A tablet and an ampule formulation containing 4-[3-(carboxymethyl)-4-[{E}-2-[4-(4-

L13 ANSWER 1 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)



Double bond geometry as shown.

PAGE 1-A



L13 ANSWER 1 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)
 phenylbutoxy)phenyl]ethenyl-1H-indol-1-yl)butanoic acid were described.
 ACCESSION NUMBER: 2006-886288 HCAPLUS
 DOCUMENT NUMBER: 145:292868
 TITLE: Preparation of indole derivatives as leukotriene receptor antagonists
 INVENTOR(S): Takeuchi, Jun; Nakayama, Yoshiyuki; Fujita, Manabu
 PATENT ASSIGNEE(S): Ono Pharmaceutical Co., Ltd., Japan
 SOURCE: PCT Int. Appl., 353pp.
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

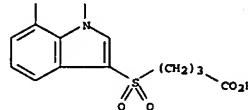
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RN: AD, BE, BG, CH, CI, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CP, CO, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TO, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, T2, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
PRIORITY APPLN. INFO.:			JP 2005-51392	A 20050225
			JP 2005-352787	A 20051207

OTHER SOURCE(S): MARPAT 145:292868
 IT 908137-43-9P 908137-44-0P
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (preparation of indole-derivs. as leukotriene receptor antagonists for prevention and/or treatment of respiratory diseases)
 RN 908137-43-9 HCAPLUS
 CN 1H-Indole-1-acetic acid, 3-[(3-carboxypropyl)sulfonyl]-7-[{1E}-2-[4-(4-phenoxybutoxy)phenyl]ethenyl]- (9CI) (CA INDEX NAME)

Double bond geometry as shown.

L13 ANSWER 1 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)

PAGE 2-A



REFERENCE COUNT: 49 THERE ARE 49 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE REFORMAT

L13 ANSWER 2 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN
 ED Entered STN: 08 Jun 2006
 GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB The invention relates to indoles and related compds. of formula I, which are modulators of peroxisome proliferator-activated receptors (PPARs).

In compds. I, U, V, W, X, Y and Z are independently selected from N and (un)substituted C; R1 is selected from (un)substituted carboxyl and carboxylic acid isosteres; R2 is H, (un)substituted lower alkyl, (un)substituted lower alkenyl, (un)substituted cycloalkyl, (un)substituted heterocyclyl, (un)substituted aryl, (un)substituted arylsulfonyl, etc.; R3 and R4 are independently selected from H, (un)substituted lower alkyl, (un)substituted cycloalkyl, (un)substituted heterocyclyl, etc., or R3 and R4 may combine to form a 3- to 7-membered (un)substituted cycloalkyl or 3- to 7-membered (un)substituted heterocyclyl; and n is 0, 1, or 2; provided that no more than two of U, V,

W, and Y are N. The invention also relates to the preparation of I, pharmaceutical compns. comprising a compound of formula I and a pharmaceutically acceptable carrier, as well as to the use of the compns. for the treatment or prevention of a disease or condition for which PPAR modulation provides a therapeutic benefit. Arylsulfonylation of 6-methoxyindole with 4-tolylthiol and oxidation gave sulfonylindole II, which underwent alkylation with Me 3-bromopropionate and ester hydrolysis to give indolepropionic acid III. Some compds. of the invention express

EC50 values below 1 nM for at least one PPAR subtype and some compds. exhibit at least 5-fold selectivity for one subtype over other subtypes of receptors (no specific data).

ACCESSION NUMBER: 2006:528077 HCAPLUS

DOCUMENT NUMBER: 145:45938

TITLE: Indoles and related derivatives as PPAR modulators, their preparation, pharmaceutical compositions, and use in therapy.

INVENTOR(S): Lin, Jack; Ibrahim, Prabha N.; Artis, Dean R.; Zhang, Chao; Wang, Weiru; Shi, Shenghua

PATENT ASSIGNEE(S): Plexikon, Inc.- USA

SOURCE: PCT Int. Appl., 95 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

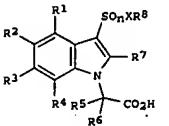
LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006060535	A2	20060608	WO 2005-US43412	20051129

L13 ANSWER 3 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN
 ED Entered STN: 09 May 2005
 GI



AB Title compds. (I; R1-R4 = H, halo, alkyl, alkoxy, OH, NO2, cyano, N(R9)2, SO2R9, CON(R9)2, etc.; R9 = H, alkyl; R5, R6 = H, alkyl; RSR6 = C3-7 cycloalkyl; R7 = H, alkyl; n = 1, 2; Z = bond, NR9; R8 = (substituted) alkyl, alkenyl, alkynyl, biphenyl, (bi- or tricyclic) heteroaryl, Ph, naphthyl; with provisos), were prepared for the treatment of allergic diseases such as asthma, allergic rhinitis and atopic dermatitis. Thus, (5-fluoro-2-methyl-3-phenylmethane)sulfonyl-1H-indol-1-yl)acetic acid (general preparation given) showed CRTH2 binding with Ki = 6 nM.

ACCESSION NUMBER: 2005:395269 HCAPLUS

DOCUMENT NUMBER: 142:463597

TITLE: Preparation of indol-1-yacetates as chemoattractant receptor-homologous molecule expressed on TH2 cells (CRTH2) antagonists

INVENTOR(S): Middlemiss, David; Ashton, Mark Richard; Boyd, Edward Andrew; Brookfield, Frederick Arthur; Armer, Richard Edward

PATENT ASSIGNEE(S): Oxagen Limited, UK

SOURCE: PCT Int. Appl., 49 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005040114	A1	20050506	WO 2004-GB4336	20040103

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AU 2004283139 A1 20050506 AU 2004-283139 20041013
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 EP 1675826 A1 20060705 EP 2004-768867 20041013
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L13 ANSWER 2 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)

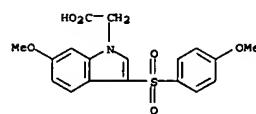
WO 2006060535 A3 20060914
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US 2006135540 A1 20060622 US 2005-289781 20051129
 PRIORITY APPLN. INFO.: US 2004-631893P P 20041130

US 2005-715250P P 20050907

OTHER SOURCE(S): MARPAT 145:45938
 IT 889129-65-1P, 2-(6-Methoxy-3-((4-methoxybenzene)sulfonyl)indol-1-yl)acetic acid
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (drug candidate; preparation of indoles and related compds. as PPAR modulators)

RN 889129-65-1 HCAPLUS
 CN 1H-Indole-1-acetic acid, 6-methoxy-3-[(4-methoxyphenyl)sulfonyl]- (9CI) (CA INDEX NAME)



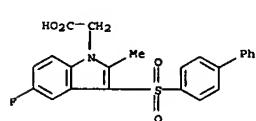
L13 ANSWER 3 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)

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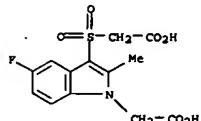
GB 2004-216 A 20040114
 WO 2004-GB4336 W 20041013

OTHER SOURCE(S): MARPAT 142:463597
 IT 851460-57-6P 851460-58-7P 851460-59-8P
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 851460-63-4P 851460-64-5P 851460-65-6P
 851460-66-7P 851460-67-8P 851460-68-9P
 851460-69-0P 851460-70-3P 851460-71-4P
 851460-72-5P 851460-73-6P 851460-74-7P
 851460-75-8P 851460-76-9P 851460-77-0P
 851460-78-1P 851460-79-2P 851460-80-5P
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (claimed compound; preparation of indolylacetates as CRTH2 antagonists)

RN 851460-57-6 HCAPLUS
 CN 1H-Indole-1-acetic acid, 3-[(1,1'-biphenyl)-4-ylsulfonyl]-5-fluoro-2-methyl- (9CI) (CA INDEX NAME)



RN 851460-58-7 HCAPLUS
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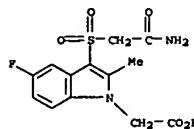


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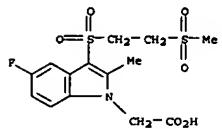
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L13 ANSWER 3 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN

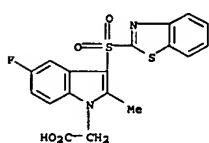
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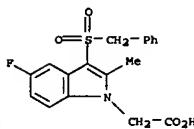


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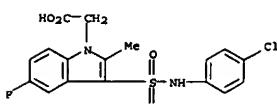


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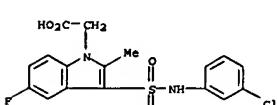
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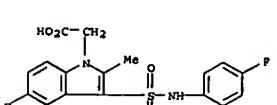
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CN 1H-Indole-1-acetic acid, 3-[(4-chlorophenyl)amino]sulfonyl]-5-fluoro-2-methyl- (9CI) (CA INDEX NAME)



RN 851460-67-8 HCAPLUS
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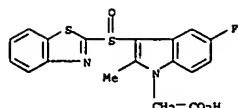


RN 851460-68-9 HCAPLUS
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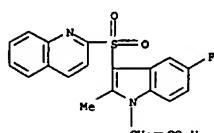


RN 851460-69-0 HCAPLUS
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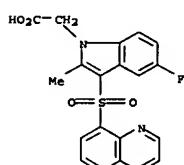
L13 ANSWER 3 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)



RN 851460-63-4 HCAPLUS
CN 1H-Indole-1-acetic acid, 5-fluoro-2-methyl-3-(2-quinolinylsulfonyl)- (9CI) (CA INDEX NAME)

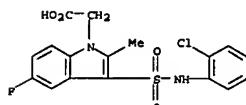


RN 851460-64-5 HCAPLUS
CN 1H-Indole-1-acetic acid, 5-fluoro-2-methyl-3-(8-quinolinylsulfonyl)- (9CI) (CA INDEX NAME)

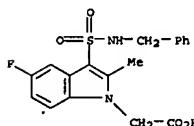


RN 851460-65-6 HCAPLUS
CN 1H-Indole-1-acetic acid, 5-fluoro-2-methyl-3-[(phenylmethyl)sulfonyl]- (9CI) (CA INDEX NAME)

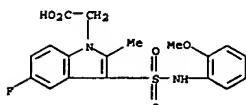
L13 ANSWER 3 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)



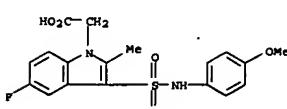
RN 851460-70-3 HCAPLUS
CN 1H-Indole-1-acetic acid, 5-fluoro-2-methyl-3-[(phenylmethyl)amino]sulfonyl]- (9CI) (CA INDEX NAME)



RN 851460-71-4 HCAPLUS
CN 1H-Indole-1-acetic acid, 5-fluoro-3-[(2-methoxyphenyl)amino]sulfonyl]-2-methyl- (9CI) (CA INDEX NAME)



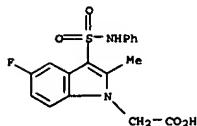
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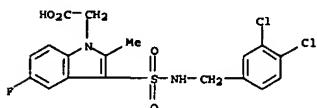
Young, Shawquia, Page 6

L13 ANSWER 3 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)

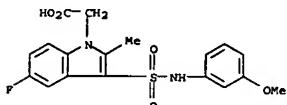
RN 851460-73-6 HCAPLUS
 CN 1H-Indole-1-acetic acid, 5-fluoro-2-methyl-3-[(phenylamino)sulfonyl]- (9CI) (CA INDEX NAME)



RN 851460-74-7 HCAPLUS
 CN 1H-Indole-1-acetic acid, 3-[[[(3,4-dichlorophenyl)methyl]amino]sulfonyl]-5-fluoro-2-methyl- (9CI) (CA INDEX NAME)

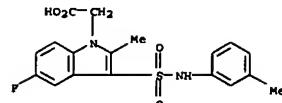


RN 851460-75-8 HCAPLUS
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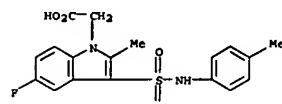


RN 851460-76-9 HCAPLUS
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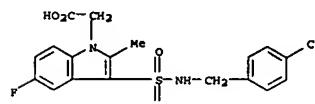
L13 ANSWER 3 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)



RN 851460-77-0 HCAPLUS
 CN 1H-Indole-1-acetic acid, 5-fluoro-2-methyl-3-[(4-methylphenyl)amino]sulfonyl]- (9CI) (CA INDEX NAME)

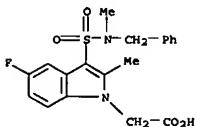


RN 851460-78-1 HCAPLUS
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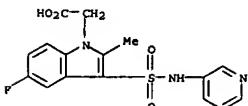


RN 851460-79-2 HCAPLUS
 CN 1H-Indole-1-acetic acid, 5-fluoro-2-methyl-3-[(methyl(phenylmethyl)amino)sulfonyl]- (9CI) (CA INDEX NAME)

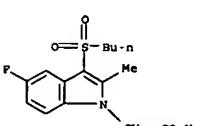
L13 ANSWER 3 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)



RN 851460-80-5 HCAPLUS
 CN 1H-Indole-1-acetic acid, 5-fluoro-2-methyl-3-[(3-pyridinylamino)sulfonyl]- (9CI) (CA INDEX NAME)



IT 851460-91-8P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (preparation of indolylacetates as CRTH2 antagonists)
 RN 851460-91-8 HCAPLUS
 CN 1H-Indole-1-acetic acid, 3-(butylsulfonyl)-5-fluoro-2-methyl- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE REFORMAT

L13 ANSWER 4 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN

ED Entered STN: 26 Jan 2004

GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB The invention relates to substituted indoles I (wherein: n = 1, 2; R1 = halogen, CN, NO2, SO2R4, OR4, SR4, S(O)R4, (hetero)aryl, (un)substituted alk(en)ynyl, etc.; R2 = H, halogen, CN, SO2R4, C(O)R4, (un)substituted alkyl, etc.; R3 = (hetero)aryl optionally containing N, S, or O, (un)substituted with halogen, CN, NO2, SO2R4, OR, OR4, SR4, S(O)R4, (un)substituted alk(en)ynyl, etc.; R4 = (hetero)aryl or alkyl (un)substituted by halogen, (hetero)aryl, etc.) as modulators of CRTH2 receptor activity useful in treatment of diseases (such as respiratory disorders) which are caused by excessive or unregulated production of PGD2.

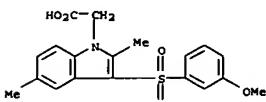
For instance, compound II (example 2, pIC50 = 8.1) was prepared via heterocyclization of 4-ClC6H4NNH2 with 4-ClC6H4SCH2C(O)CH3, transformation of obtained indole III to the Me N-acetate analog of III, subsequent N-oxidation and hydrolysis of ester.

ACCESSION NUMBER: 2004:60469 HCAPLUS
 DOCUMENT NUMBER: 140:111276
 TITLE: Preparation of indole-3-sulphur derivatives and their use in the treatment of respiratory disorders
 INVENTOR(S): Bonnert, Roger; Dickinson, Mark; Rasul, Rukhsana; Sangane, Hitesh; Teague, Simon
 PATENT ASSIGNEE(S): AstraZeneca AB, Swed.
 SOURCE: PCT Int. Appl. S9 pp.
 DOCUMENT TYPE: PCT
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004007451	A1	20040122	WO 2003-SE1216	20030715
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MM, MZ, SD, SL, SZ, T2, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CP, CO, CI, CM, GA, GN, GO, GW, ML, MR, NE, SN, TD, TG				
CA 2492445	AA	20040122	CA 2003-2492445	20030715
AU 2003251260	A1	20040202	AU 2003-251260	20030715
BR 2003012729	A	20050510	BR 2003-12729	20030715
EP 1551802	A1	20050713	EP 2003-764279	20030715
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
CN 1678579	A	20051005	CN 2003-819971	20030715
JP 2005537265	T2	20051208	JP 2004-521364	20030715
NZ 537606	A	20060831	NZ 2003-537606	20030715
ZA 2005000170	A	20060222	ZA 2005-170	20050107

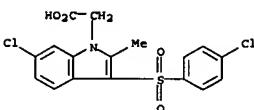
L13 ANSWER 4 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)
 NO 2005000828 A 20050405 NO 2005-828 20050216
 US 2006111426 A1 20060525 US 2005-52125 20050815
 PRIORITY APPLN. INFO.: SE 2002-2241 A 20020717
 SE 2002-3713 A 20021213
 WO 2003-SE1216 W 20030715

OTHER SOURCE(S): MARPAT 140:111276
 IT 646514-88-7P, 3-[(3-Methoxyphenyl)thio]-2,5-dimethyl-1H-indole-1-acetic acid
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (intermediate; preparation of indole-3-sulfur derivs. as modulators of CRTH2 receptor activity useful as drugs for treatment of respiratory disorders)
 RN 646514-88-7 HCAPLUS
 CN 1H-Indole-1-acetic acid, 3-[(3-methoxyphenyl)sulfonyl]-2,5-dimethyl- (9CI)
 (CA INDEX NAME)

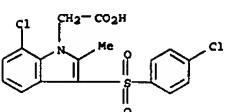


IT 646514-29-6P 646514-32-1P, 5-Chloro-3-[(4-chlorophenyl)sulfonyl]-2-methyl-1H-indole-1-acetic acid
 646514-37-6P, 6-Chloro-3-[(4-chlorophenyl)sulfonyl]-2-methyl-1H-indole-1-acetic acid 646514-42-3P, 7-Chloro-3-[(4-chlorophenyl)sulfonyl]-2-methyl-1H-indole-1-acetic acid
 646514-47-8P, 5-Chloro-3-[(4-chlorophenyl)sulfonyl]-4-cyano-2-methyl-1H-indole-1-acetic acid 646514-52-5P, 5-Chloro-3-[(4-chlorophenyl)sulfonyl]-6-cyano-2-methyl-1H-indole-1-acetic acid 646514-56-9P, 3-[(4-Chlorophenyl)sulfonyl]-2,5-dimethyl-1H-indole-1-acetic acid 646514-60-5P, 3-[(4-Chlorophenyl)sulfonyl]-4-(ethylsulfonyl)-7-methoxy-2-methyl-1H-indole-1-acetic acid 646514-67-2P, 3-[(4-Chlorophenyl)sulfonyl]-5-cyano-2-methyl-1H-indole-1-acetic acid 646514-73-0P, 3-[(4-Chlorophenyl)sulfonyl]-5-cyano-2-methyl-1H-indole-1-acetic acid 646514-79-6P, 4-Chloro-3-[(4-chlorophenyl)sulfonyl]-2-methyl-1H-indole-1-acetic acid 646514-81-2P, 3-[(2-Chlorophenyl)sulfonyl]-2,5-dimethyl-1H-indol-1-acetic acid 646514-98-9P, 3-[(4-Cyano phenyl)sulfonyl]-2,5-dimethyl-1H-indole-1-acetic acid 646515-01-7P, 3-[(2-Methylphenyl)sulfonyl]-2,5-dimethyl-1H-indol-1-acetic acid 646515-05-1P, 3-[(2-Ethylphenyl)sulfonyl]-2,5-dimethyl-1H-indol-1-acetic acid 646515-09-5P,

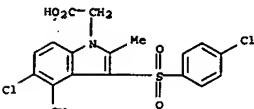
L13 ANSWER 4 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)
 RN 646514-37-6 HCAPLUS
 CN 1H-Indole-1-acetic acid, 6-chloro-3-[(4-chlorophenyl)sulfonyl]-2-methyl- (9CI) (CA INDEX NAME)



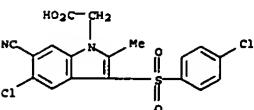
RN 646514-42-3 HCAPLUS
 CN 1H-Indole-1-acetic acid, 7-chloro-3-[(4-chlorophenyl)sulfonyl]-2-methyl- (9CI) (CA INDEX NAME)



RN 646514-47-8 HCAPLUS
 CN 1H-Indole-1-acetic acid, 5-chloro-3-[(4-chlorophenyl)sulfonyl]-4-cyano-2-methyl- (9CI) (CA INDEX NAME)



RN 646514-52-5 HCAPLUS
 CN 1H-Indole-1-acetic acid, 5-chloro-3-[(4-chlorophenyl)sulfonyl]-6-cyano-2-methyl- (9CI) (CA INDEX NAME)



L13 ANSWER 4 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)
 3-[(4-Chlorophenyl)sulfonyl]-2-methyl-4-nitro-1H-indole-1-acetic acid 646515-15-3P, 4-(Acetylaminoo)-3-[(4-chlorophenyl)sulfonyl]-2-methyl-1H-indole-1-acetic acid 646515-23-3P,

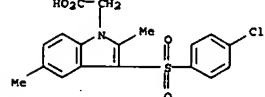
3-[(4-Chlorophenyl)sulfonyl]-2-methyl-4-[(methylsulfonyl)amino]-1H-indole-1-acetic acid 646515-31-3P, 3-[(4-Chlorophenyl)sulfonyl]-4-[(ethylamino)-2-methyl-1H-indole-1-acetic acid 646515-33-5P, 3-[(2,6-Dichlorophenyl)sulfonyl]-2,5-dimethyl-1H-indole-1-acetic acid 646515-42-6P, 3-[(4-Chlorophenyl)sulfonyl]-5-fluoro-2-methyl-1H-indole-1-acetic acid ammonium salt 646515-50-6P, 3-[(3-Chlorophenyl)sulfonyl]-5-fluoro-2-methyl-1H-indole-1-acetic acid ammonium salt 646515-54-0P, 5-Fluoro-2-methyl-3-[(4-(trifluoromethyl)phenyl)sulfonyl]-1H-indole-1-acetic acid ammonium salt 646515-59-5P, 3-[(4-Chlorophenyl)-5-fluoro-2-methyl-1H-indole-1-acetic acid 646515-61-9P, 3-[(3-Chlorophenyl)sulfonyl]-5-fluoro-2-methyl-1H-indole-1-acetic acid 646515-63-1P, 5-Fluoro-2-methyl-3-[(4-(trifluoromethyl)phenyl)sulfonyl]-1H-indole-1-acetic acid 646515-66-6P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepns. of indole-3-sulfur derivs. as modulators of CRTH2 receptor activity useful as drugs for treatment of respiratory disorders)

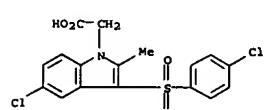
RN 646514-29-6 HCAPLUS

CN 1H-Indole-1-acetic acid, 3-[(4-chlorophenyl)sulfonyl]-2,5-dimethyl- (9CI) (CA INDEX NAME)



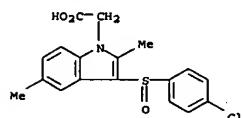
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CN 1H-Indole-1-acetic acid, 5-chloro-3-[(4-chlorophenyl)sulfonyl]-2-methyl- (9CI) (CA INDEX NAME)

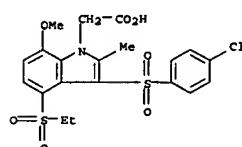


L13 ANSWER 4 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)

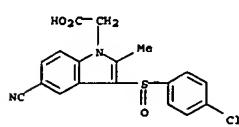
RN 646514-56-9 HCAPLUS
 CN 1H-Indole-1-acetic acid, 3-[(4-chlorophenyl)sulfonyl]-2,5-dimethyl- (9CI) (CA INDEX NAME)



RN 646514-60-5 HCAPLUS
 CN 1H-Indole-1-acetic acid, 3-[(4-chlorophenyl)sulfonyl]-4-(ethylsulfonyl)-7-methoxy-2-methyl- (9CI) (CA INDEX NAME)



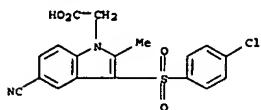
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 CN 1H-Indole-1-acetic acid, 3-[(4-chlorophenyl)sulfonyl]-5-cyano-2-methyl- (9CI) (CA INDEX NAME)



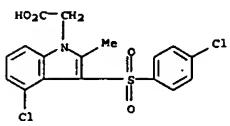
RN 646514-73-0 HCAPLUS
 CN 1H-Indole-1-acetic acid, 3-[(4-chlorophenyl)sulfonyl]-5-cyano-2-methyl- (9CI) (CA INDEX NAME)

L13 ANSWER 4 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN

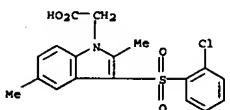
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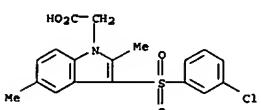
RN 646514-79-6 HCAPLUS
CN 1H-Indole-1-acetic acid, 4-chloro-3-[(4-chlorophenyl)sulfonyl]-2-methyl- (9CI) (CA INDEX NAME)



RN 646514-91-2 HCAPLUS
CN 1H-Indole-1-acetic acid, 3-[(2-chlorophenyl)sulfonyl]-2,5-dimethyl- (9CI) (CA INDEX NAME)



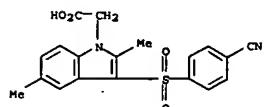
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CN 1H-Indole-1-acetic acid, 3-[(3-chlorophenyl)sulfonyl]-2,5-dimethyl- (9CI) (CA INDEX NAME)



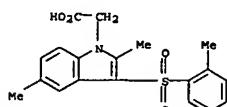
L13 ANSWER 4 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)

L13 ANSWER 4 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)

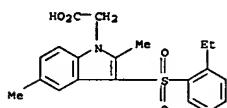
RN 646514-98-9 HCAPLUS
CN 1H-Indole-1-acetic acid, 3-[(4-cyanophenyl)sulfonyl]-2,5-dimethyl- (9CI) (CA INDEX NAME)



RN 646515-01-7 HCAPLUS
CN 1H-Indole-1-acetic acid, 2,5-dimethyl-3-[(2-methylphenyl)sulfonyl]- (9CI) (CA INDEX NAME)



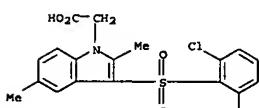
RN 646515-05-1 HCAPLUS
CN 1H-Indole-1-acetic acid, 3-[(2-ethylphenyl)sulfonyl]-2,5-dimethyl- (9CI) (CA INDEX NAME)



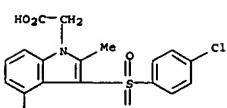
RN 646515-09-5 HCAPLUS
CN 1H-Indole-1-acetic acid, 3-[(4-chlorophenyl)sulfonyl]-2-methyl-4-nitro- (9CI) (CA INDEX NAME)

L13 ANSWER 4 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)

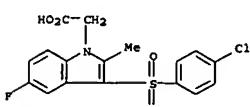
L13 ANSWER 4 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)
CN 1H-Indole-1-acetic acid, 3-[(2,6-dichlorophenyl)sulfonyl]-2,5-dimethyl- (9CI) (CA INDEX NAME)



RN 646515-36-8 HCAPLUS
CN 1H-Indole-1-acetic acid, 3-[(4-chlorophenyl)sulfonyl]-2-methyl-4-phenyl- (9CI) (CA INDEX NAME)

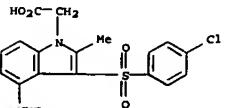


RN 646515-42-6 HCAPLUS
CN 1H-Indole-1-acetic acid, 3-[(4-chlorophenyl)sulfonyl]-5-fluoro-2-methyl-, ammonium salt (9CI) (CA INDEX NAME)



RN 646515-50-6 HCAPLUS
CN 1H-Indole-1-acetic acid, 3-[(3-chlorophenyl)sulfonyl]-5-fluoro-2-methyl-, ammonium salt (9CI) (CA INDEX NAME)

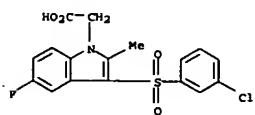
RN 646515-31-3 HCAPLUS
CN 1H-Indole-1-acetic acid, 3-[(4-chlorophenyl)sulfonyl]-4-(ethylamino)-2-methyl- (9CI) (CA INDEX NAME)



RN 646515-33-5 HCAPLUS

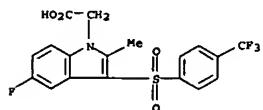
L13 ANSWER 4 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN

(Continued)

● NH₃

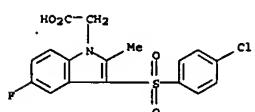
RN 646515-54-0 HCAPLUS

CN 1H-Indole-1-acetic acid, 5-fluoro-2-methyl-3-[(4-(trifluoromethyl)phenyl)sulfonyl]-, ammonium salt (9CI) (CA INDEX NAME)

● NH₃

RN 646515-59-5 HCAPLUS

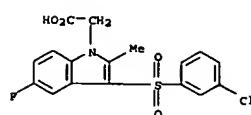
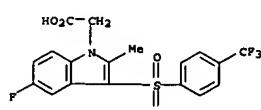
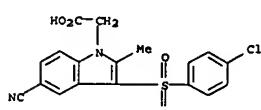
CN 1H-Indole-1-acetic acid, 3-[(4-chlorophenyl)sulfonyl]-5-fluoro-2-methyl- (9CI) (CA INDEX NAME)



RN 646515-61-9 HCAPLUS

CN 1H-Indole-1-acetic acid, 3-[(3-chlorophenyl)sulfonyl]-5-fluoro-2-methyl- (9CI) (CA INDEX NAME)

L13 ANSWER 4 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN (Continued)

RN 646515-63-1 HCAPLUS
CN 1H-Indole-1-acetic acid, 5-fluoro-2-methyl-3-[(4-(trifluoromethyl)phenyl)sulfonyl]- (9CI) (CA INDEX NAME)RN 646518-01-6 HCAPLUS
CN 1H-Indole-1-acetic acid, 3-[(4-chlorophenyl)sulfonyl]-5-cyano-2-methyl-, sodium salt (9CI) (CA INDEX NAME)

● Na

REFERENCE COUNT:

3

THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT